

Office use only: IDNUM:
KIDCODE:



# NATIONAL CENSUS OF INFANTS, CHILDREN & YOUTH WITH DEAFBLINDNESS IN NORTH DAKOTA:

*Instructions:* Please check and update all information for accuracy. Also, complete any missing information. Return form to:

Sherri Nelson, Coordinator North Dakota Dual Sensory Project 500 Stanford Road Grand Forks, ND 58203

#701-231-6033 Note, information on this page is not included in the Census Report. Child's Name: First Name Middle Initial Last Name Date of Birth: \_\_\_\_\_ Female Male Child's Home Special Education Unit: Phone number: \_\_\_\_\_ School/ Agency Serving Child: Address: \_\_\_\_\_ Parent/ Guardian Name: Address: \_\_\_\_ Work Phone: \_\_\_\_ Home Phone: Does the child reside on an Indian Reservation? Yes If yes, which one. No Fort Berthold Indian Reservation..... Turtle Mountain Band of Chippewa Indian Reservation Standing Rock Sioux Tribe Indian Reservation Spirit Lake Indian Reservation (Devils Lake Sioux Tribe) **PERSON COMPLETING THIS FORM:** (Please fill out completely) Name/Title: \_\_\_\_\_\_ Phone Number: \_\_\_\_\_ Address: \_\_\_\_\_\_

Agency/School: E-mail Address:



# Child Count Code Sheet 12-1-2010

Column 4 - Gender	
0. Male	1. Female

Column 8 - Primary Identified Etiology		
Hereditary/Chromosomal Syndromes and Disorders		
101 Aicardi syndrome	130 Marshall syndrome	
102 Alport syndrome	131 Maroteaux-Lamy syndrome (MPS VI)	
103 Alstrom syndrome	132 Moebius syndrome	
104 Apert syndrome (Acrocephalosyndactyly, Type 1)	133 Monosomy 10p	
105 Bardet-Biedl syndrome (Laurence Moon-Biedl)	134 Morquio syndrome (MPS IV-B)	
106 Batten disease	135 NF1 - Neurofibromatosis (von Recklinghausen disease)	
107 CHARGE association	136 NF2 - Bilateral Acoustic Neurofibromatosis	
108 Chromosome 18, Ring 18	137 Norrie disease	
109 Cockayne syndrome	138 Optico-Cochleo-Dentate Degeneration	
110 Cogan Syndrome	139 Pfieffer syndrome	
111 Cornelia de Lange	140 Prader-Willi	
112 Cri du chat syndrome (Chromosome 5p- syndrome)	141 Pierre-Robin syndrome	
113 Crigler-Najjar syndrome	142 Refsum syndrome	
114 Crouzon syndrome (Craniofacial Dysotosis)	143 Scheie syndrome (MPS I-S)	
115 Dandy Walker syndrome	144 Smith-Lemli-Opitz (SLO) syndrome	
116 Down syndrome (Trisomy 21 syndrome)	145 Stickler syndrome	
117 Goldenhar syndrome	146 Sturge-Weber syndrome	
118 Hand-Schuller-Christian (Histiocytosis X)	147 Treacher Collins syndrome	
119 Hallgren syndrome	148 Trisomy 13 (Trisomy 13-15, Patau syndrome)	
120 Herpes-Zoster (or Hunt)	149 Trisomy 18 (Edwards syndrome)	
121 Hunter Syndrome (MPS II)	150 Turner syndrome	
122 Hurler syndrome (MPS I-H)	151 Usher I syndrome	
123 Kearns-Sayre syndrome	152 Usher II syndrome	
124 Klippel-Feil sequence	153 Usher III syndrome	
125 Klippel-Trenaunay-Weber syndrome	154 Vogt-Koyanagi-Harada syndrome	
126 Kniest Dysplasia	155 Waardenburg syndrome	
127 Leber congenital amaurosis	156 Wildervanck syndrome	
128 Leigh Disease	157 Wolf-Hirschhorn syndrome (Trisomy 4p)	
129 Marfan syndrome	199 Other	
<b>Pre-Natal/Congenital Complications</b>	Post-Natal/Non-Congenital Complications	
201 Congenital Rubella	301 Asphyxia	
202 Congenital Syphilis	302 Direct Trauma to the eye and/or ear	
203 Congenital Toxoplasmosis	303 Encephalitis	
204 Cytomegalovirus (CMV)	304 Infections	
205 Fetal Alcohol syndrome	305 Meningitis	
206 Hydrocephaly	306 Severe Head Injury	
207 Maternal Drug Use	307 Stroke	
208 Microcephaly	308 Tumors	
209 Neonatal Herpes Simplex (HSV)	309 Chemically Induced	
299 Other	399 Other	
Related to Prematurity	Undiagnosed	
401 Complications of Prematurity	501 No Determination of Etiology	

Column 9 - Race/Ethnicity		_
<ol> <li>American Indian or Alaska</li> <li>Asian</li> <li>Black or African American</li> <li>Hispanic/Latino</li> </ol>	Native	<ul><li>5. White</li><li>6. Native Hawaiian/Pacific Islander</li><li>7. Two or more races</li></ul>
Column 10 – Documented Vision L	.oss (Items 5 and 8 are intent	tionally not used and they are unavailable as an option)
<ol> <li>Low Vision</li> <li>Legally Blind</li> <li>Light Perception Only</li> <li>Totally Blind</li> </ol>	,	<ul> <li>6. Diagnosed Progressive Loss</li> <li>7. Further Testing Needed (1 year only)</li> <li>9. Documented Functional Vision Loss</li> </ul>
Column 11 -Cortical Vision Impair	rment	
0. No	1. Yes	2. Unknown
<ol> <li>Mild</li> <li>Moderate</li> <li>Moderately Severe</li> <li>Severe</li> </ol>		<ol> <li>Profound</li> <li>Diagnosed Progressive Loss</li> <li>Further Testing Needed (1 year only)</li> <li>Documented Functional Hearing Loss</li> </ol>
Column 13 - Central Auditory Pro	cessing Disorder	
0. No	1. Yes	2. Unknown
Column 14 - Auditory Neuropathy		
0. No	1. Yes	2. Unknown
Column 15 - Cochlear Implant		·
0. No	1. Yes	2. Unknown
Column 16 - Other Impairments of	or Conditions-Orthopedic/I	Physical
0. No		1. Yes

0. No	1. Yes	
Column 18- Other Impairments or	Conditions-Behavioral	
0. No	1. Yes	
	1	
Column 19- Other Impairments or Conditions-Complex Health Care Needs		
0. No	1. Yes	
Column 20- Other Impairments or	Conditions-Communication, Speech/Language	
0. No	1. Yes	
Column 21- Other Impairments or	Conditions	
0. No	1. Yes	

Column 23 - Part C Category Code	
<ol> <li>At-risk for developmental delays (as defined by the state's Part C Lead Agency)</li> <li>Developmentally Delayed</li> </ol>	Also included for Child Count reporting purposes are: 888. Not Reported Under Part C

Column 22 is intentionally not used. (Previously this column was titled "Funding Category".)

Column 24 - Part B Category Code		
1. Mental Retardation	9. Deaf-blindness	
2. Hearing Impairment (includes deafness)	10. Multiple Disabilities	
3. Speech or Language Impairment	11. Autism	
4. Visual Impairment (includes blindness)	12. Traumatic Brain Injury	
5. Emotional Disturbance	13. Developmentally Delayed-age 3 through 9	
6. Orthopedic Impairment	Also included for Child Count reporting purposes are:	
7. Other Health Impairment	14. Non-Categorical	
8. Specific Learning Disability	888. Not Reported under Part B of IDEA	

Column 25 - Early Intervention Setting (Birth through 2)		
1. Home	2. Community-based settings	3. Other settings

### Column 26 - Educational Setting (3-21)

#### ECSE (3-5) Settings

- 1. Attending a regular early childhood program at least 80% of the time
- Attending a regular early childhood program 40% to 79% of the time
- 3. Attending a regular early childhood program less than 40 % of the time
- 4. Attending a separate class
- 5. Attending a separate school
- 6. Attending a residential facility
- 7. Service provider location
- 8. Home

#### School aged (6-21) settings

- 9. Inside the regular class 80% or more of day
- 10. Inside the regular class 40% to 79% of day
- 11. Inside the regular class less than 40% of day
- 12. Separate school
- 13. Residential facility
- 14. Homebound/Hospital
- 15. Correctional facilities
- 16. Parentally placed in private schools

## Column 27 - Participation in Statewide Assessments

- 1. Regular grade-level state assessment
- 2. Regular grade-level state assessment with accommodations
- 3. Alternate assessments aligned with gradelevel achievement standards
- 4. Alternate assessments based on alternate achievement standards
- 5. Modified achievement standards
- 6. Not yet required

## **Column 28 - Part C Exiting Status (Birth through 2)**

- 0. In a Part C early intervention program
- 1. Completion of IFSP prior to reaching maximum age for Part C
- 2. Eligible for IDEA, Part B
- 3. Not eligible for Part B, exit with referrals to other programs
- 4. Not eligible for Part B, exit with no referrals

- 5. Part B eligibility not determined
- 6. Deceased
- 7. Moved out of state
- 8. Withdrawal by parent (or guardian)
- 9. Attempts to contact the parent and/or child were unsuccessful

### Column 29 - Part B Exiting

- 0. In ECSE or school-aged special education program
- 1. Transferred to regular education
- 2. Graduated with regular diploma
- 3. Received a certificate
- 4. Reached maximum age

- 5. Died
- 6. Moved, known to be continuing
- 7. (intentionally not used)
- 8. Dropped out

Column 30 – Deaf-Blind Project Exiting Status	
Eligible to receive services from the deaf-blind project	No longer eligible to receive services from the state deaf-blind project

Column 31 - Living Setting	
1. Home: With parents	6. Group home (less than 6 residents)
2. Home: Extended family	7. Group home (6 or more residents)
3. Home: Foster parents	8. Apartment (with non-family person(s))
4. State residential facility	9. Pediatric nursing home
5. Private residential facility	555. Other (Specify)

Column 32 - Corrective Lenses		
0. No	1. Yes	2. Unknown

Column 33 - Assistive Li	istening Devices	
0. No	1. Yes	2. Unknown

Column 34 - Additional Assistive Technology			
0. No	1. Yes		2. Unknown

# **Submittal Instructions**

# All data are due by May 1st for inclusion in the National Deaf-Blind Child Count Summary.

Please call or email Mark Schalock or Robbin Bull for any additional information or clarifications related to the Child Count reporting process, or with any NCDB FMP database questions.

Please e-mail a copy of your completed report to Mark Schalock.

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